

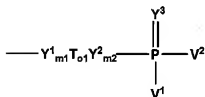
AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Claim 1 (Currently amended): A coated stent for implantation in human vessels, orifices, and conduits, and for creating and sustaining openings there and for preventing restenosis thereof after implantation, wherein the coated stent comprises ~~comprising~~ a stent structure coated with a compound containing a high density, negatively charged domain of at least three vicinally oriented phosphorus-containing radicals.

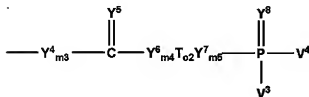
Claim 2 (Currently amended): A coated stent according to claim 1 wherein the phosphorus-containing radicals have the following formula:

a)



or

b)



wherein

V^1 to V^4 are $\text{Y}^8_{m6}\text{T}_{o3}\text{U}$

T_{o1} to T_{o3} are $(\text{CH}_2)_n$, CHCH , or $\text{CH}_2\text{CHCHCH}_2$

o1 to o3 are 0 to 1

n is 0 to 4

U is $R^1Y^9_{m7}$, $CY^{10}Y^{11}R^2$, $SY^{12}Y^{13}Y^{14}R^3$,

$PY^{15}Y^{16}Y^{17}R^4R^5$,

$Y^{18}PY^{19}Y^{20}Y^{21}R^6R^7$, CH_2NO_2 , $NHSO_2R^8$ or

$NHCY^{22}Y^{23}R^9$

m1 to m7 are 0 to 1

Y^1 to Y^{23} are NR^{10} , NOR^{11} , O or S

and where R^1 to R^{11} are

i) hydrogen;

ii) a straight or branched saturated or unsaturated alkyl residue containing 1-22 carbon atoms;

iii) a saturated or unsaturated aromatic or non-aromatic homo- or heterocyclic residue containing 3-22 carbon atoms and 0-5 heteroatoms consisting of nitrogen, oxygen or sulfur;

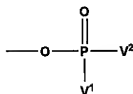
iv) a straight or branched saturated or unsaturated alkyl residue containing 1-22 carbon atoms substituted with a saturated or unsaturated aromatic or non-aromatic homo- or heterocyclic residue containing 3-22 carbon atoms and 0-5 heteroatoms consisting of nitrogen, oxygen or sulfur; and

v) an aromatic or non-aromatic homo- or heterocyclic residue containing 3-22 carbon atoms and 0-5 heteroatoms consisting of nitrogen, oxygen or sulfur substituted with a straight or branched saturated or unsaturated alkyl residue containing 1-22 carbon atoms[.].

in the said groups ii-v, the residues and/or the ~~substituents~~ substituents thereof being substituted with 0-6 of the following groups: hydroxy, alkoxy, aryloxy, acyloxy, carboxy, alkoxycarbonyl, alkoxycarbonyloxy, aryloxy carbonyl, aryloxy carbonyloxy,

carbamoyl, fluoro, chloro, bromo, azido, cyano, oxo, oxa, amino, imino, alkylamino, arylamino, acylamino, arylazo, nitro, alkylthio or alkylsulfonyl.

Claim 3 (Currently amended): A coated stent according to claim 2 wherein the phosphorus-containing radicals have the following formula:



wherein V^1 and V^2 are OH, $(\text{CH}_2)_p\text{OH}$, COOH, CONH₂, CONOH, $(\text{CH}_2)_p\text{COOH}$, $(\text{CH}_2)_p\text{CONH}_2$, $(\text{CH}_2)_p\text{CONOH}$, $(\text{CH}_2)_p\text{SO}_3\text{H}$, $(\text{CH}_2)_p\text{SO}_3\text{NH}_2$, $(\text{CH}_2)_p\text{NO}_2$, $(\text{CH}_2)_p\text{PO}_3\text{H}_2$, $\text{O}(\text{CH}_2)_p\text{OH}$, $\text{O}(\text{CH}_2)_p\text{COOH}$, $\text{O}(\text{CH}_2)_p\text{CONH}_2$, $\text{O}(\text{CH}_2)_p\text{CONOH}$, $(\text{CH}_2)_p\text{SO}_3\text{H}$, $\text{O}(\text{CH}_2)_p\text{SO}_3\text{NH}_2$, $\text{O}(\text{CH}_2)_p\text{NO}_2$, $\text{O}(\text{CH}_2)_p\text{PO}_3\text{H}_2$, CF_2COOH and p is 1 to 4.

Claim 4 (Original): A coated stent according to claim 3 wherein the phosphorus-containing radicals are phosphate groups.

Claim 5 (Currently amended): A coated stent according to ~~anyone~~ any one of claims 1-4 wherein a backbone to the high density negatively charged region of vicinally oriented phosphorus-containing radicals is a cyclic moiety.

Claim 6 (Original): A coated stent according to claim 5 wherein the backbone is a saturated or unsaturated aromatic or non-aromatic homo- or heterocyclic moiety where the heteroatom is nitrogen, oxygen, sulfur or selenium.

Claim 7 (Original): A coated stent according to claim 6 wherein the cyclic moiety comprises 4 to 24 atoms, preferably 5 to 18 atoms.

Claim 8 (Original): A coated stent according to claim 7 wherein the cyclic moiety is selected from the group of cyclopentane, cyclohexane, cycloheptane, inositol, monosaccharide,

disaccharide, trisaccharide, tetrasaccharide, piperidin, tetrahydrothiopyran, 5-oxotetrahydrothiopyran, 5,5-dioxotetrahydrothiopyran, tetrahydroselenophyran, tetrahydrofuran, pyrrolidine, tetrahydrothiophene, 5-oxotetrahydrothiophene, 5,5-dioxotetrahydrothiophene, tetrahydroselenophene, benzene, cumene, mesitylene, naphthalene and phenanthrene.

Claim 9 (Currently amended): A coated stent according to claim 8 ~~wherein~~ wherein the cyclic moiety is selected from the group of alloinositol, cisinositol, epiinositol, D/L-chiroinositol, scylloinositol, myoinositol, mucinositol and neoinositol.

Claim 10 (Currently amended): ~~The use~~ The coated stent according to claim 8 wherein the cyclic moiety is selected from the group of D/L-ribose, D/L-arabinose, D/L-xylose, D/L-lyxose, D/L-allose, D/L-altrose, D/L-glucose, D/L-mannose, D/L-gulose, D/L-idose, D/L-galactose, D/L-talose, D/L-ribulose, D/L-xylulose, D/L-psicose, D/L-sorbose, D/L-tagatose and D/L-fructose.

Claim 11 (Original): A coated stent according to claim 3 wherein one of the phosphorus-containing radicals is axial and, two of the phosphorus-containing radicals are equatorial.

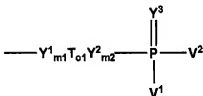
Claim 12 (Original): A coated stent according to claim 11 wherein the compound is selected from the group of myo-inositol-1,2,6-trisphosphate, myo-inositol-hexa-kis-phosphate, mannose-2,3,4-trisphosphate, rhamnose-2,3,4-trisphosphate, galactose-2,3,4-trisphosphate, methyl-6-O-butyl- α -D-mannopyranoside-2,3,4-trisphosphate, 1,5-anhydro-D-arabinitol-2,3,4-trisphosphate, fructose-2,3,4-trisphosphate, 1,2-O-ethylene- β -D-fructopyranoside-2,3,4-trisphosphate, cyclohexane-1,2,3-triol trisphosphate, 1,5-dideoxy-1,5-iminoarabinitol-2,3,4-trisphosphate, altrose-2,3,4-trisphosphate, methyl-6-O-butyl- α -D-altropyranoside-2,3,4-trisphosphate or derivatives thereof.

Claim 13 (Currently amended): A method of selecting a ~~The use~~ a restenosis ~~resistant~~ resistant stent for implantation to in a human patient; the method comprising the steps of:

selecting a coated stent for implantation in human vessels, orifices, and conduits, and for creating and sustaining openings there and for preventing, alleviating or ~~combating~~ combating restenosis thereof after implantation, wherein the coated stent comprises ~~comprising~~ a stent structure coated with a compound containing a high density, negatively charged domain of at least three vicinally oriented phosphorus-containing radicals.

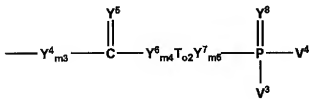
Claim 14 The ~~use~~ method according to claim 13 wherein the compound containing phosphorus-containing radicals have the following formula:

a)

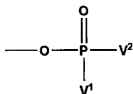


or

b)



Claim 15 (Currently amended): The ~~use~~ method according to claim 13 where the compound containing phosphorus-containing radicals have the following formula:



Claim 16 (Currently amended): The method according to claim 13 wherein the compound is selected from the group of myo-inositol-1,2,6-trisphosphate, myo-inositol-hexakis-phosphate, mannose-2,3,4-trisphosphate, rhamnose-2,3,4-trisphosphate, galactose-2,3,4-trisphosphate, methyl-6-O-butyl- α -D-mannopyranoside-2,3,4-trisphosphate, 1,5-anhydro-D-arabinitol-2,3,4 trisphosphate, fructose-2,3,4-trisphosphate, 1,2-O-ethylene- β -D-fructopyranoside-2,3,4-trisphosphate, cyclohexane-1,2,3-triol trisphosphate, 1,5-dideoxy-1,5-iminoarabinitol-2,3,4-trisphosphate, altrose-2,3,4-trisphosphate, methyl-6-O-butyl- α -D-altrpyranoside-2,3,4-trisphosphate or derivatives thereof.